Attorney Docket No. 27353-501 UTIL

## **AMENDMENT**

## Amendments to the Claims:

Please amend the claims as follows, without prejudice:

## In the Claims:

- 1. (Currently Amended) A method of generating a protein array from <u>a library</u> a plurality of two or more non-identical target DNA sequences, the method comprising:
- (a) inserting a marker DNA sequence in frame immediately following a start codon of each of the plurality of target DNA sequences or immediately preceding a stop codon of each of a the plurality of target DNA sequences or both, to form a plurality of a library of modified DNA sequences which encode a plurality a library of modified amino acid sequences each comprising a marker moiety;
- (b) expressing the plurality <u>library</u> of modified <u>target</u> amino acid sequences from the <u>plurality library</u> of modified DNA sequences;
- (c) purifying and immobilizing each of the <del>plurality of modified target</del> amino acid sequences to a solid support in a single step, wherein the marker moietyies of the <del>plurality of target</del> modified amino acid sequences is <u>are directly attached</u> to the <u>single solid support</u> in a spatially defined format, thereby generating a protein array, and
  - (d) washing said solid support to remove unbound proteins.
- 2. (Previously Presented) The method as claimed in claim 1 wherein the marker moiety is a peptide sequence selected from the group consisting of:
  - (a) a histidine tag;
  - (b) a complete protein or protein domain;
  - (c) a maltose binding protein domain;
  - (d) an antibody epitope;
  - (e) biotin or a biotin mimic;
  - (f) a glutathione-S-transferase (GST) domain; and
  - (g) a peptide sequence which effects attachment to the solid support.

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3. (Previously Presented) The method as claimed in claim 1 wherein the marker moiety

allows for purification of the individual proteins in the array.

4. (Previously Presented) The method of claim 1 wherein the marker DNA sequence is

inserted such that the start or stop codon for each of the proteins is replaced.

5-7. (Canceled).

8. (Withdrawn) A method of screening one or more compounds for biological activity

which comprises:

(a) bringing said one or more compounds into contact with the array made according

to any one of claims 1 to 4; and

(b) measuring binding of the one or more compounds to the proteins in the array.

9. (Withdrawn) A method of screening one or more proteins for specific protein-protein

interactions which comprises the step of bringing said one or more proteins into contact with an

array made according to any one of claims 1 to 4, and measuring binding of the one or more

specific proteins with the proteins of the array.

10. (Withdrawn) A method of screening one or more proteins for specific interactions with

one or more nucleic acid probes which comprises the step of bringing said one or more nucleic

acid probes into contact with an array made according to any one of claims 1 to 4, and measuring

binding of the probes to the proteins in the array.

11. (Withdrawn) A method for the rapid screening of a test compound, test protein or test

nucleic acid, the method comprising:

(a) contacting the test compound, test protein or test nucleic acid with a spatially

defined array produced according to any one of claims 1-4 comprising a plurality of array bound

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proteins, with each array bound protein being at a different position on a solid support, wherein

the plurality of array bound proteins comprises a plurality of different proteins expressed in a

single species; and

(b) detecting any interaction between the array bound proteins and the test compound,

test protein or test nucleic acid.

12. (Withdrawn) A method of screening for molecules which recognize each protein in the

array, the method comprising:

(a) contacting the molecules with a spatially defined array comprising a plurality of

array bound proteins produced according to any one of claims 1-4, with each array bound protein

being at a different position on a solid support, wherein the plurality of array bound proteins

comprises a plurality of different proteins expressed in a single species; and

(b) detecting any interaction between the array bound proteins and the molecules.

13. (Previously Presented) A method of generating an antibody array which comprises

(a) bringing a protein array, made according to any one of claims 1 to 4, into contact

with an antibody library, such that one or more proteins in the protein array bind to at least one

antibody in the antibody library;

(b) removing any unbound antibodies; and

(c) immobilisation of those antibodies bound to proteins in the protein array.

14. (Withdrawn) A method for the screening of protein function or abundance which

comprises the step of bringing an antibody array as defined in claim 13 into contact with a

mixture of one or more proteins.

15. (Canceled).

16. (Cancelled)

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17. (Cancelled)

18. (Previously Presented) The method of any one of claims 1 to 4 wherein the protein array

comprises serine proteases, kinases or p450 enzymes.

19. (Currently Amended) The method of any one of claims 1 to 4 wherein said plurality

<u>library</u> of modified amino acid sequences are modified human amino acid sequences.

20. (Previously Presented) The method of claim 2 wherein the marker moiety is selected

from the group consisting of FLAG and Strep.

21. (Previously Presented) The method of claim 1 or 2 wherein the marker moiety is post-

translationally modified.

22. (Previously Presented) The method of claim 21 wherein the post-translational

modification comprises the addition of a biotin or a lipid molecule.

23. (Previously Presented) The method of claim 1 wherein said modified amino acid

sequences are folded into the correct conformation.

24. (Currently Amended) The method of claim 1 wherein said inserting step inserts a marker

DNA sequence in frame immediately following a start codon of each plurality target DNA

sequence and immediately preceding a stop codon of each plurality target DNA sequence, to form

a plurality library of modified DNA sequences which encode a plurality library of modified

amino acid sequences each comprising two marker moieties.

25. (Withdrawn) A method of generating a proteomic array of proteins of unknown amino

acid sequences comprising the steps of:

(a) providing a cDNA library as a plurality of target DNA sequences; and

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(b) generating a protein array using the method of any of claims 1 to 4 to produce a

proteomic array of proteins of unknown amino acid sequence.

26. (Currently Amended) A method of screening for antibodies which recognize each protein

in the array, the method comprising:

(a) contacting the antibodies with a spatially defined array comprising a plurality

library of array bound proteins produced according to any one of claims 1-4, with each array

bound protein being at a different position on a solid support, wherein the pluralitylibrary of

array bound proteins comprises a plurality library of different proteins expressed in a single

species; and

(b) detecting any interaction between the array bound proteins and the antibodies.

27. (Previously Presented) The method of claim 1, wherein the marker moiety provides a

high-affinity attachment to the solid support.

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